

## CLAIMS

### WHAT IS CLAIMED IS:

- 1        1.        A method for screening a plurality of test substances useful for the  
2        prevention or treatment of a disease involving an oxidative stress, which comprises  
3        the steps of
  - 4                i)    testing each of the test substances for its ability to inhibit the activity of  
5                GADD34L and
  - 6                ii)   identifying the test substance which inhibits the activity of GADD34L,  
7                thereby to identify a test substance useful as a preventive or therapeutic agent for a  
8                disease involving an oxidative stress.
- 1        2.        A method for identifying a test substance useful for the prevention or  
2        treatment of a disease involving an oxidative stress, which comprises testing a test  
3        substance for its ability to inhibit the activity of GADD34L, thereby to determine  
4        whether the substance promotes resistance to cell stress, and to identify said  
5        substance as a preventive or therapeutic agent for a disease involving an oxidative  
6        stress.
- 1        3.        The method according to claim 1 or 2, wherein the test substance inhibits  
2        the activity of the GADD34L protein by disrupting formation of the GADD34L and  
3        PP1c protein complex.
- 1        4.        The method according to claim 1 or 2, wherein the test substance inhibits  
2        the activity of GADD34L by inhibiting the production of GADD34L protein from the  
3        GADD34L mRNA.
- 1        5.        The method according to claim 1 or 2, wherein the test substance inhibits  
2        the activity of GADD34L by inhibiting the production of GADD34L mRNA from the  
3        GADD34L genomic locus.
- 1        6.        The method according to claim 2 or 3, further comprising a step of  
2        verifying whether said test substance does not cause stress to cells.

1        7.        The method according to claim 1 or 2, which comprises the steps of  
2                i)        contacting the test substance or each of the test substances with a cell-  
3        free composition containing GADD34L and PP1c proteins in the form of a purified  
4        complex and eIF2 $\alpha$  in a phosphorylated form,  
5                ii)        assessing the level of phosphorylation of eIF2 $\alpha$ , in comparison with  
6        the level of phosphorylation determined in the absence of test substances, in a cell-  
7        free composition containing GADD34L and PP1c proteins in the form of a purified  
8        complex and eIF2 $\alpha$  in a phosphorylated form, and  
9                iii)        identifying the test substance which provides a higher level of  
10       phosphorylation of eIF2 $\alpha$ , in comparison with the level of phosphorylation  
11       determined in the absence of test substance, thereby to identify a test substance useful  
12       as a preventive or therapeutic agent for a disease involving an oxidative stress.

1       8.        The method according to claim 7, wherein the assessment of the level of  
2       phosphorylation of eIF2 $\alpha$  is effected by an immunoassay using an antibody that  
3       specifically recognizes the phosphorylated form of eIF2 $\alpha$ .

1       9.        The method according to claim 7, wherein the assessment of the level of  
2       phosphorylation of eIF2 $\alpha$  is effected by tracking the covalent binding of a  
3       radiolabelled phosphate group to eIF2 $\alpha$ .

1       10.       The method according to claim 1 or 2, which comprises the steps of  
2                i)        contacting a test substance or each of the test substances with cells not  
3        subject to stress that contain PP1c and eIF2 $\alpha$  and that overexpress GADD34L, or  
4        portions thereof,  
5                ii)        assessing the level of phosphorylation of eIF2 $\alpha$  after contact with the test  
6        substance or test substances, in comparison with the level of eIF2 $\alpha$  phosphorylation  
7        in the absence of test substances, and  
8                iii)        identifying the test substance which provides a higher level of  
9        phosphorylation of eIF2 $\alpha$ , in comparison with the level of phosphorylation  
10       determined in the absence of test substance, thereby to identify a test substance useful  
11       as a preventive or therapeutic agent for a disease involving an oxidative stress.

1 11. The method according to claim 10, wherein the assessment of the level of  
2 phosphorylation of eIF2 $\alpha$  is effected by an immunoassay using an antibody that  
3 specifically recognizes the phosphorylated form of eIF2 $\alpha$ .

1 12. The method according to claim 10, wherein the assessment of the level of  
2 phosphorylation of eIF2 $\alpha$  is effected by tracking the covalent binding of a  
3 radiolabelled phosphate group to eIF2 $\alpha$ .

1 13. The method according to claim 1 or 2, which comprises the steps of,  
2 i) contacting a test substance or each of the test substances with cells that  
3 normally express endogenous GADD34L,

4 ii) and identifying a test substance that inhibits the expression of endogenous  
5 GADD34L, thereby to identify a test substance useful as a preventive or therapeutic  
6 agent for a disease involving an oxidative stress.

1 14. The method according to claim 13, wherein the level of GADD34L  
2 expression is assessed by determining the level of transcription of GADD34L.

1 15. The method according to claim 14, wherein determination of the level of  
2 transcription of GADD34L is effected by means of a Northern blot.

1 16. The method according to claim 14, wherein determination of the level of  
2 transcription of GADD34L is effected by means of *in situ* hybridization.

1 17. The method according to claim 13, wherein the level of GADD34L  
2 expression is assessed by the level of translation of GADD34L.

1 18. The method according to claim 17, wherein determination of the level of  
2 translation of GADD34L is effected by means of an immunoassay.

1 19. The method according to claim 1 or 2, which comprises the steps of  
2 i) contacting a test substance or each of the test substances with cells not  
3 subject to stress that overexpress GADD34L, or portions thereof,

4           ii) assessing the expression of expression of a target gene, and  
5           iii) identifying a test substance that activates the expression of the target gene,  
6 thereby to identify a test substance useful as a preventive or therapeutic agent for a  
7 disease involving an oxidative stress.

1       20.       The method according to claim 19, where the target gene is the CHOP  
2 gene.

1       21.       The method according to claim 1 or 2, which comprises the steps of,  
2           i) obtaining cells not subject to stress that overexpress GADD34L, or  
3 portions thereof, and have been transfected with a reporter gene operatively  
4 associated with all or part of the promoter of a target gene,  
5           ii) contacting a test substance or each of the test substances with these cells,  
6 and assaying the level of expression of said reporter gene, and  
7           iii) identifying a test substance that activates the expression of the reporter  
8 gene, thereby to identify a test substance useful as a preventive or therapeutic agent  
9 for a disease involving an oxidative stress.

1       22.       The method according to claim 21, where the target gene is the CHOP  
2 gene.

1       23.       The method according to claim 21, wherein said reporter gene encodes  
2 one of the group consisting of GFP, CAT, GAL, LUC, and GUS.

1       24.       The method according to claim 1 or 2, which comprises the steps of,  
2           i) obtaining cells not subject to stress that overexpress GADD34L, or  
3 portions thereof,  
4           ii) contacting a test substance or each of the test substances with the cells, in  
5 the presence of a toxic agent that induces oxidative stress,  
6           iii) quantitating cell survival of the cells that overexpress GADD34L, or  
7 portions of GADD34L, following exposure to the toxic agent in the presence and  
8 absence of test substances, and

9           iv) identifying a test substance that promotes cell survival of the cells  
10 following exposure to concentrations of toxic agent that induce oxidative stress,  
11 thereby to identify a test substance useful as a preventive or therapeutic agent for a  
12 disease involving an oxidative stress.

1       25.       The method according to claim 24 wherein the toxic agent which induces  
2 oxidative stress is tunicamycin, arsenite, or glutamate.

1       26.       The method according to claim 1 or 2, wherein the identified test  
2 substance is useful for the prevention or treatment of a disease involving neuronal  
3 ischemia.

1       27.       The method according to claim 1 or 2, wherein the identified test  
2 substance is useful for the prevention or treatment of a disease involving heart  
3 ischemia.

1       28.       The method according to claim 1 or 2, wherein the identified test  
2 substance is useful for the prevention or treatment of renal damage induced by  
3 ischemia or toxins.

1       29.       The method according to claim 1 or 2, wherein the identified test  
2 substance is useful for the prevention or treatment of an auto-immune disease.

1       30.       The method according to claim 1 or 2, wherein the selected compound is  
2 useful for the prevention or treatment of a neurodegenerative disorder.

1       31.       A method for the prevention or treatment of a disease involving an  
2 oxidative stress in a patient in need of such treatment, which comprises administering  
3 to the patient an effective amount of a GADD34L inhibitor identified for its ability to  
4 promote resistance to cell stress while not causing stress.

1       32.       A method of claim 31, wherein the disease is a disease involving neuronal  
2 ischemia, a disease involving heart ischemia, a disease involving renal damage  
3 induced by ischemia or toxins, an auto-immune disease, or a neurodegenerative  
4 disorder.